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APPENDIX O TO PART 121—HAZARDOUS MATE-RIALS TRAINING REQUIREMENTS FOR CER-TIFICATE HOLDERS

APPENDIX P TO PART 121—REQUIREMENTS FOR ETOPS AND POLAR OPERATIONS

AUTHORITY: 49 U.S.C. 106(f), 106(g), 40113, 40119, 41706, 44101, 44701-44702, 44705, 44709-44711, 44713, 44716-44717, 44722, 46105.2.

EFFECTIVE DATE NOTE: By Amdt. 121-366, 78 FR 67836, Nov. 12, 2013, the authority citation to part 121 was revised, effective Mar. 12, 2014. For the convenience of the user, the revised text is set forth as follows:

AUTHORITY: 49 U.S.C. 106(f), 106(g), 40113, 40119, 41706, 44101, 44701-44702, 44705, 44709-44711, 44713, 44716-44717, 44722, 46105; Pub. L. 111-216, 124 Stat. 2348 (49 U.S.C. 44701 note).

### SPECIAL FEDERAL AVIATION REGULATION No. 50-2

EDITORIAL NOTE: For the text of SFAR No. 50-2, see part 91 of this chapter.

SPECIAL FEDERAL AVIATION REGULATION No. 71

EDITORIAL NOTE: For the text of SFAR No. 71, see part 91 of this chapter.

#### SPECIAL FEDERAL AVIATION REGULATION No 97

EDITORIAL NOTE: For the text of SFAR No. 97, see part 91 of this chapter.

SPECIAL FEDERAL AVIATION REGULATION No. 106—Rules for use of port-ABLE OXYGEN CONCENTRATOR SYS-TEMS ON BOARD AIRCRAFT

Section 1. Applicability—This rule prescribes special operating rules for the use of portable oxygen concentrator units on board civil aircraft. This rule applies to both the aircraft operator and the passenger using the portable oxygen concentrator on board the aircraft.

Section 2. Definitions-For the purposes of this SFAR the following definitions apply: Portable Oxygen Concentrator: means the AirSep FreeStyle, AirSep LifeStyle, AirSep Focus, AirSep FreeStyle 5, Delphi RS-00400, DeVilbiss Healthcare iGo, Inogen One, Inogen One G2, Inogen One G3, Inova Labs LifeChoice, Inova Labs LifeChoice Activox,

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LifeChoice. International Biophysics Invacare XPO2, Invacare Solo2, Oxlife Independence Oxygen Concentrator, Oxus RS-Precision Medical EasyPulse. Respironics EverGo, Respironics SimplyGo. SeQual Eclipse and SeQual SAROS Portable Oxygen Concentrator medical device units as long as those medical device units: (1) Do not contain hazardous materials as determined by the Pipeline and Hazardous Materials Safety Administration; (2) are also regulated by the Food and Drug Administration; and (3) assist a user of medical oxygen under a doctor's care. These units perform by separating oxygen from nitrogen and other gases contained in ambient air and dispensing it in concentrated form to the user.

Section 3. Operating Requirements-

- (a) No person may use and no aircraft operator may allow the use of any portable oxygen concentrator device, except the AirSep FreeStyle, AirSep LifeStyle, AirSep Focus, AirSep FreeStyle 5, Delphi RS-00400, DeVilbiss Healthcare iGo, Inogen One, Inogen One G2, Inogen One G3, Inova Labs LifeChoice, Inova Labs LifeChoice Activox, International Biophysics LifeChoice. Invacare XPO2, Invacare Solo2, Oxlife Independence Oxygen Concentrator, Oxus RS-EasyPulse. Precision Medical Respironics EverGo, Respironics SimplyGo, SeQual Eclipse and SeQual SAROS Portable Oxygen Concentrator units. These units may be carried on and used by a passenger on board an aircraft provided the aircraft operator ensures that the following conditions are satisfied:
- (1) The device does not cause interference with the electrical, navigation or communication equipment on the aircraft on which the device is to be used;
- (2) No smoking or open flame is permitted within 10 feet of any seat row where a person is using a portable oxygen concentrator.
- (3) During movement on the surface, takeoff, and landing, the unit must:
- (i) Either be stowed under the seat in front of the user, or in another approved stowage location, so that it does not block the aisle way or the entryway into the row; or
- (ii) If it is to be operated by the user, be used only at a seat location that does not restrict any passenger's access to, or use of, any required emergency or regular exit, or the aisle(s) in the passenger compartment;
- (4) No person using a portable oxygen concentrator is permitted to sit in an exit row;
- (5) The pilot in command must be apprised whenever a passenger brings and intends to use a portable oxygen concentrator on board the aircraft and the pilot in command must be informed about the contents of the physician's written statement (as required in Section 3(b)(3) of this SFAR), including the magnitude and nature of the passenger's oxygen needs.

- (6) Whenever the pilot in command turns off the "Fasten Seat Belt" sign, or otherwise signifies that permission is granted to move about the passenger cabin, passengers operating their portable oxygen concentrator may continue to operate it while moving about the cabin.
- (b) The user of the portable oxygen concentrator must comply with the following conditions to use the device on board the aircraft:
- (1) The user must be capable of hearing the unit's alarms, seeing the alarm light indicators, and have the cognitive ability to take the appropriate action in response to the various caution and warning alarms and alarm light indicators, or be travelling with someone who is capable of performing those functions:
- (2) The user must ensure that the portable oxygen concentrator is free of oil, grease or other petroleum products and is in good condition free from damage or other signs of excessive wear or abuse:
- (3) The user must inform the aircraft operator that he or she intends to use a portable oxygen concentrator on board the aircraft and must allow the crew of the aircraft to review the contents of the physician's statement. The user must have a written statement, to be kept in that person's possession, signed by a licensed physician that:
- (i) States whether the user of the device has the physical and cognitive ability to see, hear, and understand the device's aural and visual cautions and warnings and is able, without assistance, to take the appropriate action in response to those cautions and warnings;
- (ii) States whether or not oxygen use is medically necessary for all or a portion of the duration of the trip; and
- (iii) Specifies the maximum oxygen flow rate corresponding to the pressure in the cabin of the aircraft under normal operating conditions
- (4) Only lotions or salves that are oxygen approved may be used by persons using the portable oxygen concentrator device;
- (5) The user, whose physician statement specifies the duration of oxygen use, must obtain from the aircraft operator, or by other means, the duration of the planned flight. The user must carry on the flight a sufficient number of batteries to power the device for the duration of the oxygen use specified in the user's physician statement, including a conservative estimate of any unanticipated delays; and
- (6) The user must ensure that all portable oxygen concentrator batteries carried on-board the aircraft in carry-on baggage are protected from short circuit and are packaged in a manner that protects them from physical damage. Batteries protected from short circuit include: (1) Those designed with

recessed battery terminals; or (2) those packaged so that the battery terminals do not contact metal objects (including the battery terminals of other batteries). When a battery-powered oxygen concentrator is carried onboard aircraft as carry-on baggage and is not intended to be used during the flight, the battery must be removed and packaged separately unless the concentrator contains at least two effective protective features to prevent accidental operation during transport.

Section 4. Expiration Date—This SFAR No. 106 will remain in effect until further notice.

[Doc. No. FAA-2004-18596, 70 FR 40164, July 12, 2005, as amended at 71 FR 53956, Sept. 12, 2006; 74 FR 2354, Jan. 15, 2009; 75 FR 742, Jan. 6, 2010; 75 FR 39632, July 12, 2010; Amdt. 121-358, 77 FR 4220, Jan. 27, 2012; Amdt. 121-361, 77 FR 63221, Oct. 16, 2012]

# Subpart A—General

## §121.1 Applicability.

This part prescribes rules governing—

- (a) The domestic, flag, and supplemental operations of each person who holds or is required to hold an Air Carrier Certificate or Operating Certificate under part 119 of this chapter.
- (b) Each person employed or used by a certificate holder conducting operations under this part including maintenance, preventive maintenance, and alteration of aircraft.
- (c) Each person who applies for provisional approval of an Advanced Qualification Program curriculum, curriculum segment, or portion of a curriculum segment under SFAR No. 58 of 14 CFR part 121, and each person employed or used by an air carrier or commercial operator under this part to perform training, qualification, or evaluation functions under an Advanced Qualification Program under SFAR No. 58 of 14 CFR part 121.
- (d) Nonstop Commercial Air Tours conducted for compensation or hire in accordance with \$119.1(e)(2) of this chapter must comply with drug and alcohol requirements in \$\$121.455, 121.457, 121.458 and 121.459, and with the provisions of part 136, subpart A of this chapter by September 11, 2007. An operator who does not hold an air carrier certificate or an operating certificate is permitted to use a person who is otherwise authorized to perform aircraft maintenance or preventive maintenance duties and who is not subject to

anti-drug and alcohol misuse prevention programs to perform—

- (1) Aircraft maintenance or preventive maintenance on the operator's aircraft if the operator would otherwise be required to transport the aircraft more than 50 nautical miles further than the repair point closest to the operator's principal base of operations to obtain these services: or
- (2) Emergency repairs on the operator's aircraft if the aircraft cannot be safely operated to a location where an employee subject to FAA-approved programs can perform the repairs.
- (e) Each person who is on board an aircraft being operated under this part.
- (f) Each person who is an applicant for an Air Carrier Certificate or an Operating Certificate under part 119 of this chapter, when conducting proving tests.
- (g) This part also establishes requirements for operators to take actions to support the continued airworthiness of each airplane.

[Doc. No. 28154, 60 FR 65925, Dec. 20, 1995, as amended by Amdt. 121–328, 72 FR 6912, Feb. 13, 2007; Amdt. 121–336, 72 FR 63411, Nov. 8, 20071

### § 121.2 Compliance schedule for operators that transition to part 121; certain new entrant operators.

- (a) *Applicability*. This section applies to the following:
- (1) Each certificate holder that was issued an air carrier or operating certificate and operations specifications under the requirements of part 135 of this chapter or under SFAR No. 38–2 of 14 CFR part 121 before January 19, 1996, and that conducts scheduled passenger-carrying operations with:
- (i) Nontransport category turbopropeller powered airplanes type certificated after December 31, 1964, that have a passenger seat configuration of 10–19 seats;
- (ii) Transport category turbopropeller powered airplanes that have a passenger seat configuration of 20–30 seats; or
- (iii) Turbojet engine powered airplanes having a passenger seat configuration of 1-30 seats.
- (2) Each person who, after January 19, 1996, applies for or obtains an initial air carrier or operating certificate and